

Knight: Chapter 17

Work, Heat, & the 1st Law of
Thermodynamics
(The Specific Heat of Gases)

Last time...

- Heat needed to change temp by ΔT (*only* when $W=0$!) is..

$$Q = Mc\Delta T$$

Specific heat

or

$$Q = nC\Delta T$$

Molar specific heat

- Heat needed for a phase change...

$$Q = \pm ML_{f,v}$$

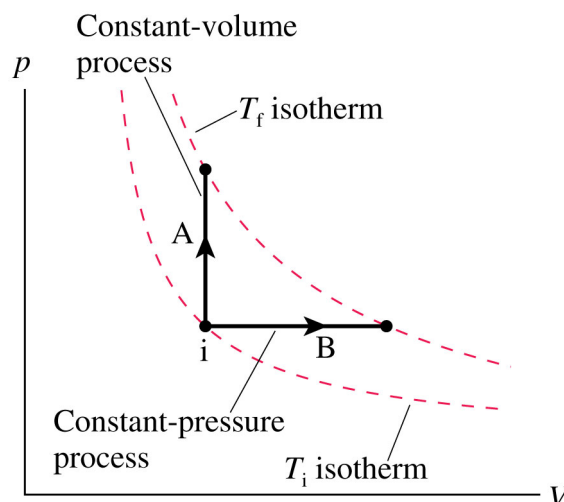
- Calorimetry...

$$Q_{net} = 0$$

The Specific Heat of Gases

Consider the two processes A & B...

- Both have the *same* ΔT , therefore the *same* ΔE_{th} , but they require *different* amounts of Q .
 - The reason is that work is done in process B but not in process A.
- The total change in thermal energy for any process, due to work and heat, is:



$$\Delta E_{th} = nC_V \Delta T \quad (\text{any ideal-gas process})$$

$$\Delta E_{th} = nC_V \Delta T = W + Q$$

The Specific Heat of Gases

Define 2 different versions of the specific heat of gases...

- one for constant-*volume* processes.
- one for constant-*pressure* processes.
- The quantity of heat needed to change the temperature of n moles of gas by ΔT is:

$$Q = nC_V\Delta T \quad (\text{temp change at constant volume})$$

$$Q = nC_P\Delta T \quad (\text{temp change at constant pressure})$$

- where C_V is the *molar specific heat at constant volume* & C_P is the *molar specific heat at constant pressure*.

Constant volume : $W=0 \therefore Q=nC_V\Delta T$ (no area under PV curve)

Constant pressure : $W \neq 0 \therefore Q=nC_P\Delta T$



The Specific Heat of Gases

What if neither the pressure nor the volume is constant?

The Specific Heat of Gases

What if neither the pressure nor the volume is constant?

- no direct way to relate Q to ΔT !
- Use the 1st law of thermodynamics:

$$Q = \Delta E_{th} - W$$

Extra credit problem

i.e. 17.8:

Calorimetry with a gas and a solid

The interior volume of a 200 g hollow aluminum box is 800 cm³. The box contains nitrogen gas at STP. A 20 cm³ block of copper at a temperature of 300° C is placed inside the box, then the box is sealed.

What is the final temperature?

$$\overset{\text{gas}}{\Delta E_{\text{in}}} = nC_V \Delta T = Q + \cancel{W} \quad \text{no work}$$

$$Q = nC_V \Delta T$$

$$m_{\text{AL}} = 0.2 \text{ kg}$$

$$Q_{\text{Cu}} + Q_{\text{AL}} + Q_{\text{N}_2} = 0$$

$$V_{\text{N}_2} = 800 \text{ cm}^3 = 8.0 \times 10^{-4} \text{ m}^3$$

$$\mu_{\text{Cu}} C_{\text{Cu}} (T_i - T_{\text{eq}}) + \mu_{\text{AL}} C_{\text{AL}} (T_i - T_{\text{eq}}) + nC_V (T_i - T_{\text{eq}}) = 0$$

$$T_{\text{eq}} = T_{\text{N}_2} = 273 \text{ K} \quad (-Q) \text{ gains heat}$$

$$(\mu_{\text{Cu}} C_{\text{Cu}} + \mu_{\text{AL}} C_{\text{AL}} + nC_V) T_i = \mu_{\text{Cu}} C_{\text{Cu}} T_{\text{eq}} + \mu_{\text{AL}} C_{\text{AL}} T_{\text{eq}} + nC_V T_{\text{eq}}$$

$$P_{\text{N}_2} = 1.01 \times 10^5 \text{ Pa}$$

$$V_{\text{Cu}} = 20 \text{ cm}^3 = 2.0 \times 10^{-5} \text{ m}^3$$

$$T_i = \frac{\mu_{\text{Cu}} C_{\text{Cu}} T_{\text{eq}} + \mu_{\text{AL}} C_{\text{AL}} T_{\text{eq}} + nC_V T_{\text{eq}}}{(\mu_{\text{Cu}} C_{\text{Cu}} + \mu_{\text{AL}} C_{\text{AL}} + nC_V)}$$

$$T_{\text{eq}} = 573 \text{ K} \quad (-Q) \text{ loses heat}$$

$$C_{\text{AL}} = 900 \text{ J/kg K}$$

$$C_{\text{Cu}} = 385 \text{ J/kg K}$$

$$C_V = 20.8 \text{ J/mol K}$$

$$T_i = 356 \text{ K}$$

$$\rho_{\text{Cu}} = \frac{\mu_{\text{Cu}}}{V_{\text{Cu}}}$$

$$\mu_{\text{Cu}} = (8920 \text{ kg/m}^3) (2.0 \times 10^{-5} \text{ m}^3)$$

$$\mu_{\text{Cu}} = 0.178 \text{ kg}$$

$$n_{\text{N}_2} = \frac{PV}{RT} = \frac{(1.01 \times 10^5 \text{ Pa})(8.0 \times 10^{-4} \text{ m}^3)}{(8.31 \text{ J/mol K})(273 \text{ K})} = 3.53 \times 10^{-2} \text{ mol}$$

C_P and C_V

Consider Table 17.4...

TABLE 17.4 Molar specific heats of gases
(J/mol K)

Gas	C_P	C_V	$C_P - C_V$
Monatomic Gases			
He	20.8	12.5	8.3
Ne	20.8	12.5	8.3
Ar	20.8	12.5	8.3
Diatomic Gases			
H ₂	28.7	20.4	8.3
N ₂	29.1	20.8	8.3
O ₂	29.2	20.9	8.3

C_P and C_V

Consider Table 17.4...

TABLE 17.4 Molar specific heats of gases (J/mol K)

Gas	C_P	C_V	$C_P - C_V$
Monatomic Gases			
He	20.8	12.5	8.3
Ne	20.8	12.5	8.3
Ar	20.8	12.5	8.3
Diatomic Gases			
H ₂	28.7	20.4	8.3
N ₂	29.1	20.8	8.3
O ₂	29.2	20.9	8.3

Notice:

- ▣ *Molar specific heats of monatomic gases are all alike.*
- ▣ *Molar specific heats of diatomic gases are very nearly alike.*
- ▣ *The difference is the same in every case &...*

$$C_p - C_V = R$$

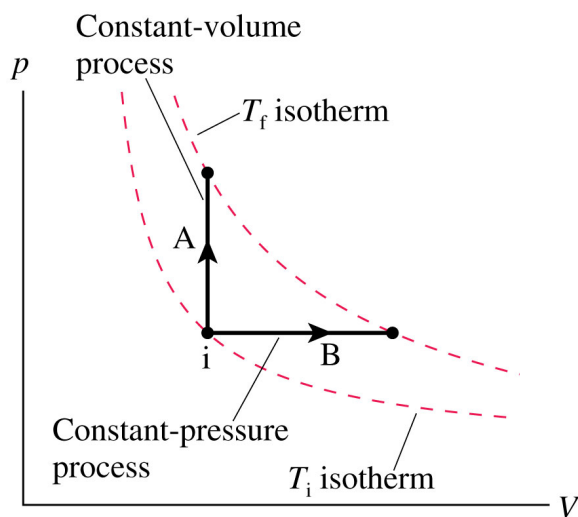
Coincidences?

C_P and C_V

The change in thermal energy of the gas is the *same* for any two processes that have the same ΔT .

$$\Delta E_{th} = nC_V \Delta T$$

Consider Figure 17.20...



Q: Which process requires *more heat* to produce the same ΔT ?

Path A

$$W=0 \therefore \Delta E_{th} = nC_V \Delta T = Q_A$$

$$\Delta E_{thB} = nC_V \Delta T = W + Q = -nRT + nC_P \Delta T \therefore C_V = -R + C_P$$

$$nC_V \Delta T = -p \Delta V + Q$$

$$R = C_P - C_V$$

$$\Delta pV = \Delta(nRT)$$

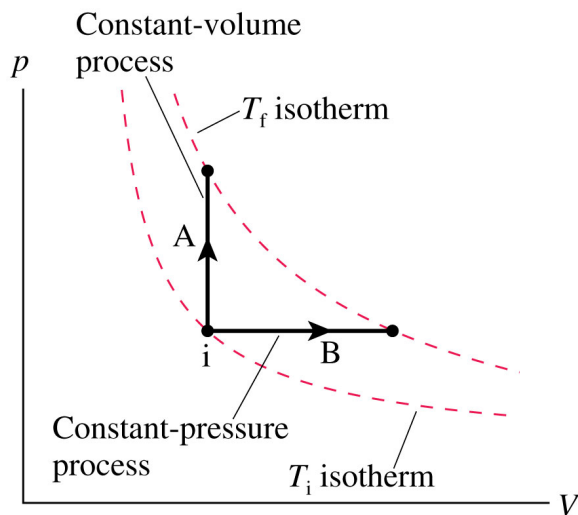
$$p \Delta V = nR \Delta T$$

C_P and C_V

The change in thermal energy of the gas is the *same* for any two processes that have the same ΔT .

$$\Delta E_{th} = nC_V \Delta T$$

Consider Figure 17.20...



$$C_P = C_V + R$$

The *heat* required to bring about a temperature change depends on what the process is.

Quiz Question 1

1 mol of air has an initial temperature of 20°C . 200 J of heat energy are transferred to the air in an isochoric process, then 200 J are removed in an isobaric process. Afterward, the air temperature is

$$C_p = C_v + R$$

$$Q = nC_p\Delta T_p \quad \text{isobaric}$$

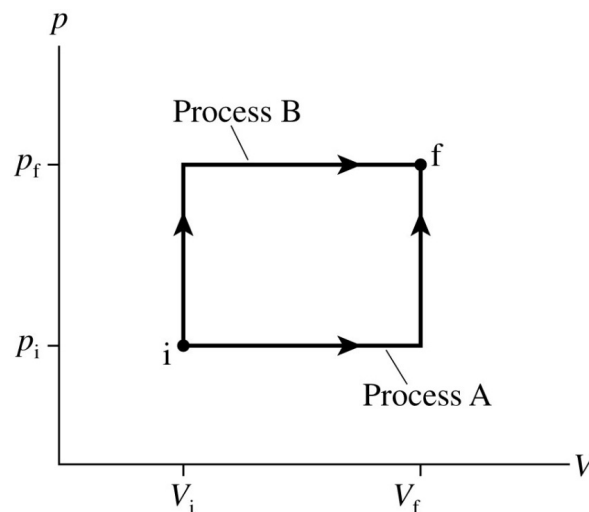
$$Q = nC_v\Delta T_v \quad \text{isochoric}$$

1. $< 20^{\circ}\text{C}$.
2. $= 20^{\circ}\text{C}$.
- ③ $> 20^{\circ}\text{C}$.
4. Not enough information is given to answer the question.

Heat depends on the path..

Consider the two ideal-gas processes...

- There's more area under the process B curve, so $|W_B| > |W_A|$.
- $W < 0$ as the gas *expands*.
- ΔE_{th} is the *same* for both processes, so $W_A + Q_A = W_B + Q_B$.
- Only true if $Q_B > Q_A$.



Heat *added* or *removed* during an ideal-gas process depends on the path followed through the pV diagram!

Adiabatic processes...

2 ways one can have an adiabatic process..

Gas cylinder can be...

1. completely surrounded by *thermal insulation*.
2. *expanded* or *compressed* very rapidly where there is no time for heat to be transferred between the gas and the environment.

Adiabatic processes...

2 ways one can have an adiabatic process..

Gas cylinder can be...

1. completely surrounded by *thermal insulation*.
2. *expanded* or *compressed* very rapidly where there is no time for heat to be transferred between the gas and the environment.

Notice:

- ▣ *Adiabatic expansions or compressions* are...
 - *fast enough* to be *adiabatic*, but *slow enough* to still be considered *quasi-static*.
- ▣ For an adiabatic process, $Q = 0$ so $\Delta E_{\text{th}} = W = nC_V\Delta T$
 - Adiabatic compression (expansion) raises (lowers) the temp of a gas.



Adiabatic processes...

An *adiabatic process* is one for which...

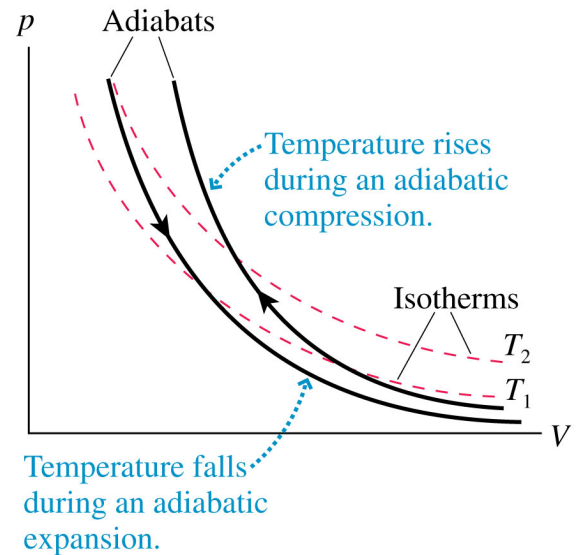
Adiabatic processes...

An *adiabatic process* is one for which...

$$pV^\gamma = \text{constant}$$

where

$$\gamma = \frac{C_P}{C_V} = 1.67 \quad \text{monatomic gas}$$
$$= 1.40 \quad \text{diatomic gas}$$



Adiabats are steeper than hyperbolic isotherms, so the temp

- ▣ *falls* during an *adiabatic expansion*
- ▣ *rises* during an *adiabatic compression*.